

AMENDMENTS TO THE CLAIMS

1-11. (Cancelled)

12. (Currently Amended) AAn *in vitro* base conversion method of a DNA sequence, which is a method of converting one or more bases in a target DNA sequence in a cell, ~~comprising~~ consisting of introducing a single-stranded DNA fragment having 300 to 3,000 bases which is prepared by cleavage from a single-stranded circular DNA, is homologous with the target DNA sequence, and contains the base(s) to be converted, into a cell, wherein the single-stranded DNA fragment is homologous with either a sense strand or an antisense strand of the target DNA.

13. (Previously Presented) The method according to claim 12, wherein the single-stranded circular DNA is a phagemid DNA.

14. (Previously Presented) The method according to claim 12, wherein the single-stranded DNA fragment is homologous with a sense strand of the target DNA sequence.

15. (Previously Presented) The method according to claim 12, wherein the target DNA sequence in the cell is a DNA sequence causing a disease due to the one or more bases.

16. (Previously Presented) The method according to claim 12, wherein one or more bases in a target DNA sequence in a cell of an organism are converted.

17. (Withdrawn) A cell in which one or more bases in a target DNA sequence have been converted by the method according to claim 12.

18. (Withdrawn) An individual organism which retains the cell according to claim 17 in the body.

19. **(Withdrawn)** A therapeutic agent, which is an agent for treating a disease caused by conversion of one or more bases in a target DNA sequence, characterized in that a single-stranded DNA fragment having 300 to 3,000 bases which is prepared from a single-stranded circular DNA, is complementary to the target DNA sequence, and contains the base(s) to be converted, has a form that can be introduced into a cell.
20. **(Withdrawn)** The therapeutic agent according to claim 19, wherein the single-stranded circular DNA is a phagemid DNA.
21. **(Withdrawn)** A therapeutic method, which is a method of treating a disease caused by conversion of one or more bases in a target DNA sequence, characterized by introducing a single-stranded DNA fragment having 300 to 3,000 bases which is prepared from a single-stranded circular DNA, is complementary to the target DNA sequence, and contains the base(s) to be converted, into a cell.
22. **(Withdrawn)** The therapeutic method according to claim 21, wherein the single-stranded circular DNA is a phagemid DNA.